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What is claimed is:

1. A method for treating or preventing vasomotor symptoms in a subject in need thereof, comprising the step of:

administering to said subject a therapeutically effective amount of at least one dual NRI/SRI compound or pharmaceutically acceptable salt thereof,

wherein said amount is less than about 37.5 mg/day.

- A method according to claim 1,
 wherein said amount is less than about 30 mg/day.
- A method according to claim 1,
 wherein said amount is less than about 25 mg/day.
- A method according to claim 1,
 wherein said amount is less than about 20 mg/day.
- A method according to claim 1,
 wherein said amount is less than about 15 mg/day.
- A method according to claim 1,
 wherein said amount is less than about 10 mg/day.
- 7. A method according to claim 1, wherein said amount is less than about 5 mg/day.
- 8. A method according to claim 1, wherein said dual NRI/SRI compound is a compound selected from the group consisting of venlafaxine, O-desmethyl-venlafaxine, milnacipran, duloxetine, and combinations and pharmaceutically acceptable salts thereof.
- 9. A method according to claim 1,

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wherein said dual NRI/SRI compound is venlafaxine or a pharmaceutically acceptable salt thereof.

10. A method according to claim 1,

wherein said dual NRI/SRI compound is O-desmethyl-venlafaxine or a pharmaceutically acceptable salt thereof.

11. A method according to claim 1,

wherein said dual NRI/SRI compound is milnacipran or a pharmaceutically acceptable salt thereof.

12. A method according to claim 1,

wherein said dual NRI/SRI compound is duloxetine or a pharmaceutically acceptable salt thereof.

13. A method according to claim 1,

wherein said vasomotor symptom is hot flush.

14. A method according to claim 1,

wherein said vasomotor symptom is caused by a thermoregulatory dysfunction.

15. A method according to claim 1,

wherein said subject is human.

16. A method according to claim 15,

wherein said human is a female.

17. A method according to claim 16,

wherein said female is pre-menopausal.

18. A method according to claim 16,

wherein said female is peri-menopausal.

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A method according to claim 16,
 wherein said female is post-menopausal.

- 20. A method according to claim 15, wherein said human is a male.
- 21. A method according to claim 20, wherein said male is naturally, chemically or surgically andropausal.